



Year: 2014

**Arthroscopic repair of traumatic isolated subscapularis tendon lesions
(Lafosse Type III or IV): a prospective magnetic resonance
imaging-controlled case series with 1 year of follow-up**

Grueninger, Patrick ; Nikolic, Nikola ; Schneider, Joerg ; Lattmann, Thomas ; Platz, Andreas ; Chmiel, Corinne ; Meier, Christoph

Abstract: **PURPOSE** The purpose of this study was to prospectively assess the efficacy of arthroscopic repair of isolated high-grade subscapularis (SSC) tendon lesions by means of clinical follow-up combined with magnetic resonance imaging investigations. **METHODS** Between January 2008 and September 2010, 11 patients (9 men and 2 women; mean age, 45 ± 10 years) with Lafosse type III or IV traumatic isolated SSC tendon lesions underwent arthroscopic repair including tenodesis of the long head of the biceps tendon. All patients were preoperatively assessed by clinical examination (Constant-Murley score [CMS]) and contrast-enhanced magnetic resonance arthrography. At 1 year of follow-up, specific clinical SSC tests, the CMS, and the loss of external rotation were evaluated. A native magnetic resonance investigation was performed to assess the structural integrity of the repair. The SSC muscle was compared with its preoperative condition regarding fatty infiltration and size (cross-sectional area). Patient satisfaction was graded from 1 (poor) to 4 (excellent). **RESULTS** The mean time interval from trauma to surgery was 3.7 months. A concomitant lesion of the biceps tendon was observed in 10 patients (91%). The mean CMS improved from 44 to 89 points ($P < .001$). The functional tests showed a significant increase in strength ($P < .05$) (belly-press test, 4.8 ± 2.9 ; lift-off test, 4.8 ± 2.9). The mean loss of external rotation at 0° of abduction was 10° compared with the contralateral side ($P < .05$). Patient satisfaction was high. Magnetic resonance imaging evaluation showed complete structural integrity of the tendon repair in all studies. The SSC showed a significant decrease in fatty infiltration and increase in the cross-sectional area. **CONCLUSIONS** Arthroscopic repair of higher-grade isolated SSC lesions provides reliable tendon healing accompanied by excellent functional results 1 year after surgery. **LEVEL OF EVIDENCE** Level IV, prospective therapeutic case series.

DOI: <https://doi.org/10.1016/j.arthro.2014.02.030>

Posted at the Zurich Open Repository and Archive, University of Zurich

ZORA URL: <https://doi.org/10.5167/uzh-98463>

Journal Article

Accepted Version

Originally published at:

Grueninger, Patrick; Nikolic, Nikola; Schneider, Joerg; Lattmann, Thomas; Platz, Andreas; Chmiel, Corinne; Meier, Christoph (2014). Arthroscopic repair of traumatic isolated subscapularis tendon lesions (Lafosse Type III or IV): a prospective magnetic resonance imaging-controlled case series with 1 year of follow-up. *Arthroscopy: The Journal of Arthroscopic Related Surgery*, 30(6):665-672.

DOI: <https://doi.org/10.1016/j.arthro.2014.02.030>

Manuscript Number: ARTH-13-784R1

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Article Type: Original Article

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Methods: Between January 2008 and September 2010 11 patients (9 male 2 female, mean age 45±10 years) with traumatic isolated SSC tendon lesions Lafosse III-IV underwent arthroscopic repair including long head of biceps tenodesis. All patients were preoperatively assessed by clinical examination (Constant-Murley score (CMS)) and contrast enhanced MR arthrography. At one year follow-up, specific clinical SSC tests, the CMS and the loss of external rotation were evaluated. A native MR investigation was performed to assess structural integrity of the repair. The SSC muscle was compared to its preoperative condition regarding fatty infiltration and size (cross-sectional area (CSA)). Patient satisfaction was graded between 4 (excellent) and 1 (poor).

Results: Mean time interval from trauma to surgery was 3.7 months. A concomitant lesion of the biceps tendon was observed in 10 patients (91%). Mean CMS improved from 44 to 89 points ($p<0.001$). The functional tests showed a significant increase of strength ($p<0.05$) (belly-press test: 4.8 vs. 2.9; lift-off test: 4.8 vs. 2.9). Mean loss of external rotation at 00 abduction was 100 compared to the contralateral side ($p<0.05$). Patient satisfaction was high. MRI evaluation showed complete structural integrity of the tendon repair in all studies. The SSC showed a significant decrease of fatty infiltration and increase of the CSA.

Conclusions: Arthroscopic repair of higher grade isolated SSC lesions provides reliable tendon healing accompanied by excellent functional results one year after surgery.

Level of Evidence: Level IV, prospective therapeutic case series.

Original Article

Arthroscopic repair of traumatic isolated subscapularis tendon lesions Lafosse III-IV: a prospective MRI-controlled case series with one year follow-up

Running title: Arthroscopic repair of subscapularis tendon lesions

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9

Abstract

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Results: Mean time interval from trauma to surgery was 3.7 months. A concomitant lesion of the biceps tendon was observed in 10 patients (91%). Mean CMS improved from 44 to 89 points ($p < 0.001$). The functional tests showed a significant increase of strength ($p < 0.05$) (belly-press test: 4.8 vs. 2.9; lift-off test: 4.8 vs. 2.9). Mean loss of external rotation at 0° abduction was 10° compared to the contralateral side ($p < 0.05$). Patient satisfaction was high. MRI evaluation showed complete structural integrity of the tendon repair in all studies. The SSC showed a significant decrease of fatty infiltration and increase of the CSA.

Conclusions: Arthroscopic repair of higher grade isolated SSC lesions provides reliable tendon healing accompanied by excellent functional results one year after surgery.

~~In our series we observed significant decrease in fatty SSC muscle infiltration and an increase of muscular mass.~~ **Level of Evidence:** ~~Therapeutic~~ Level IV, prospective therapeutic case series.

Introduction

Isolated lesions of the subscapularis (SSC) are ~~rare~~less common. Lafosse et al. reported a prevalence of 10.1% comparing isolated SSC tears with all SSC lesions ¹. The importance of the SSC regarding its biomechanical and functional properties has increasingly been ~~recognized over the last years~~. Its function includes internal rotation of the shoulder, force coupling in the transverse plane and it also contributes to the anterior stability of the shoulder. Thus, patients with impaired SSC function present with increased passive external rotation and pathological lift-off test ². Complete tears of the SSC with retraction may cause an anterior displacement of the humeral head onto the glenoid due to disruption of the force couple of the rotator cuff.

Open repair has been associated with good clinical outcomes in several studies ^{3,4}. In 2002, the technique of arthroscopic SSC repair was described by Burkhart and Tehrany ⁵. However, some of these studies also included anterosuperior lesions or did not distinguish between low-grade and higher-grade lesions ^{5,6}. Only a few study groups included follow-up imaging such as CT arthrography or MRI to assess structural integrity of the repair or muscular alterations such as fatty infiltration or muscular atrophy ^{1, 7-~~11~~10}.

The purpose of the present study was to evaluate clinical outcome after arthroscopic repair of isolated traumatic high-grade lesions of the SSC tendons. Furthermore, muscular alterations of the SSC were assessed by means of degree of fatty infiltration and muscular mass in MRI at follow-up after one year. Our hypothesis was that patients with traumatic high-grade lesion to the SSC tendon would benefit from early arthroscopic repair.

Methods

Patients. Between 01/2008 and 09/2010 eleven consecutive patients with traumatic isolated complete SSC tendon tears type III and IV according to the Lafosse classification ¹ undergoing an

all arthroscopic repair were included in ~~the study~~this prospective case series. Patients with anterosuperior lesions or mass lesions were excluded. The study was approved by the local ethics committee and informed consent for operative treatment and all the follow-up investigations was obtained from all the patients. Follow-up after one year included a clinical examination and MRI to assess the structural integrity of the repair. Fatty infiltration and the muscular mass of the SSC, SSP and ISP were also evaluated and compared to preoperative imaging.

Operative technique. All arthroscopic procedures were performed in general anesthesia with an interscalene catheter for postoperative pain control. The patient was seated in a standardized beach chair position with arm traction of 2-3 kg. A perioperative antibiotic prophylaxis with cefuroxim was routinely administered. A standard 30° arthroscope was used. According to Lafosse¹ the portals were named from A (posterior "soft spot" portal) to E (anterosuperior portal). Before introducing the arthroscope into the glenohumeral joint through the A portal, the joint was infiltrated with 20ml of diluted adrenaline (1ml of adrenaline (1mg/ml) and 19ml of normal saline) to decrease intraoperative bleeding. Furthermore, systolic blood pressure was constantly kept at a maximum of 100mmHg during surgery. With these measures the intraarticular pressure could be kept as low as 35mmHg until the end of the intervention for most patients. In some cases, the pressure had to be increased in a stepwise manner during the procedure in order to provide good visibility. However, a maximum pressure of 60mmHg was never exceeded in this series.

The diagnostic arthroscopy was performed and a probe was inserted through the D portal (anterolateral portal). After confirming the isolated lesion of the SSC, the lesion was classified according to the Lafosse classification¹. Only type III and IV lesions were included (Fig 1A and B). The long head of biceps tendon (LHBT) was inspected with a probe. For the tenodesis of the LHBT one suture of a double loaded ~~threaded suture anchor screw anchor (HEALIX 4.5mm-Ti™ Anchor w/ORTHOCORD®-violet / blue strand, Size 2, or Fastin®-RC 5mm Anchor w/ORTHOCORD®-violet / blue strand, Size 2, Mitek Sports Medicine, Raynham, USA)~~ was used performing a "lasso-loop stitch" in all patients of this series^{11,12}. The second suture of this anchor was used for the

reconstruction of the most superior part of the SSC tendon at a later stage. The rotator interval was opened using a shaver and a bipolar diathermy (VAPR® 3 Premiere90 Electrode 90° or VAPR® 3 LDS electrode 90°, Mitek Sports Medicine, Raynham, USA). The tip of the coracoid process and the conjoint tendon were dissected before the E portal (anterosuperior portal) was created under direct visualization in an outside-in technique. To achieve a 270° release of the SSC the middle glenohumeral ligament was debrided from the posterior aspect of the SSC. To facilitate mobilization, a traction suture through the D portal was routinely used for retracted SSC tendons. The superior glenohumeral ligament was also resected. A shaver and burr were used to prepare the footprint of the SSC on the lesser tuberosity. The arthroscope was now placed through the D portal and the instrumentation was changed from the D to the E portal. To complete the release of the SSC all adhesions to the coracoid were released and the subcoracoidal bursa was removed. In type IV lesions the axillary nerve was visualized routinely (Fig 2). In type III and IV lesions with no or moderate tension the reconstruction of the SSC was performed with 2 ~~FASTIN® 5mm or~~ 4.5mm HEALIX-Ti™ threaded suture anchors usually applying U-stitches. The remaining suture of the LHBT tenodesis anchor was used for a lasso-loop stitch at the upper border of the SSC achieving a pseudo double-row reconstruction (Fig. 3). In type IV lesions with higher tension (n=2) lasso-loops instead of U-stitches were used and the reconstruction was reinforced with a double-row reconstruction in suture bridge technique ~~(VERSALOK™ Anchor, Mitek Sports Medicine, Raynham, USA).~~ No coracoplasties were performed in the study group as we did not see any signs of coracoid impingement. Surgery was completed with a routine acromioplasty. However, a rather limited than a formal acromioplasty was performed in asymptomatic patients. All operations were performed by the corresponding author.

Rehabilitation. Postoperatively, all patients were immobilized on a 30° abduction pillow for six weeks. Physiotherapy was initiated on the first postoperative day starting with passive and pain free exercises for the shoulder for six weeks. During this period, external rotation was limited at

0°. Patients were encouraged to perform active wrist and elbow movements from the beginning. For personal hygiene all patients were briefed to shower with a waterproof abduction wedge. After six weeks, active exercises were started to regain full range of motion. Weight bearing and strengthening exercises were allowed after twelve weeks. Depending on the kind of sports, a gradual return to these activities was allowed not before six months postoperatively.

Clinical evaluation. Preoperatively, the patient's history in general and the trauma mechanism and the time of the injury in particular were recorded. A standardized physical examination was performed before the operation and at follow-up by the first author. The clinical examination included a Constant-Murley score (CMS) ^{42,13}. The SSC function was tested with the modified belly-press test and the modified lift-off test as described by Lafosse et. al.¹. Muscular strength was graded from 0 to 5 according to the classification of neurological assessment. External rotation in 0° abduction was measured and the loss of external rotation compared to the contralateral side was recorded. Internal rotation was assessed according to the CMS subscore for internal rotation. The loss of internal rotation was defined by the loss of points compared to the contralateral side (CMS subscore for internal rotation). At follow-up, our patients were also asked to rate their level of satisfaction ranging from poor, fair and good to excellent.

Radiological evaluation. All patients underwent a standardized radiographic evaluation including a true anteroposterior radiograph in neutral rotation and an axillary view before and immediately after surgery. Preoperatively, all patients were also evaluated with contrast enhanced Magnetic Resonance Imaging (Arthro-MRI). Fatty infiltration of the SSC was graded according to Goutallier ^{43,14}, modified by Fuchs ⁴⁴⁻¹⁵ for MRI from 0 to 4. Accordingly, grade 0 indicates no fatty infiltration; grade 1, some fatty streaks; grade 2, less fat than muscle; grade 3, as much fat as muscle; and grade 4, more fat than muscle. The cross-sectional area (CSA) of the SSC was measured according to the method proposed by Zanetti et al. ⁴⁵⁻¹⁶ employing standard measurement tools in our PACS software. The CSA was measured in mm² at the most lateral

image on which the scapular spine was in contact with the rest of the scapula in the sagittal reconstructions.

At follow-up after one year, an MRI investigation with a dedicated shoulder-coil (Magnetom Avanto 1.5T, Siemens Medical Solution, Erlangen, Germany) without contrast enhancement was obtained. On the axial and paracoronal views the tendons of the supraspinatus (SSP), infraspinatus (ISP) and SSC were evaluated regarding continuity and retraction. Tendon integrity was assessed in axial and sagittal T2-weighted and proton density-weighted sequences (Fig. 4). A tendon re-rupture was diagnosed if a clear retraction was present or if a gap in a tendon was filled with a water equivalent signal. The fatty muscular infiltration and CSA of the SSP, ISP and SSC were determined as described above (Fig. 5). The integrity of the tenodesis of the LHBT was evaluated on the most superior axial cross-section where the bicipital groove was still visible. In contrast to the preoperative MR arthrographies only native MRI studies were performed at follow-up. Acceptance of an invasive technique for pure scientific reasons without direct benefit to the patient may be low and may also be discussed controversially by ethical aspects. Furthermore, the superiority of MR arthrography for evaluating structural integrity of SSC repair is not proven¹⁷. The application of intraarticular gadolinium does not change the appearance of the muscles such as the SSC on the images. The complete pre- and postoperative radiological assessment was performed by one experienced MR-radiologist with special training in musculoskeletal imaging. This radiologist ~~who~~ was blinded to the clinical results.

Statistical analysis. Results were analyzed using statistical software (IBM SPSS version 20). Statistical analysis was performed using the paired t-test and the Wilcoxon signed rank test when appropriate. All data are presented as means with standard deviations (ranges are provided in brackets). A double sided P-value of <0.05 was considered statistically significant.

Results

Between 01/2008 and 09/2010 a total of 11 traumatic isolated SSC tendon lesions Lafosse III/IV were included into our prospective consecutive case series. Mean age of the patients was 45 ± 10 years (range 32-65), 9 males and 2 females. The dominant shoulder was affected in eight patients (82%). In 5 patients, a forceful external rotation was reported, 3 patients fell on their outstretched arm, and 2 patients suffered from a first episode of a traumatic anterior shoulder displacement. In 1 patient, the trauma mechanism could not be clarified.

Mean interval from trauma to surgery was 3.7 ± 4.7 months (range 0.3-13.3 months). All patients had a full clinical follow-up investigation after 1 year. However, one patient who presented with an excellent clinical outcome, refused MRI due to ~~ageraphobia~~claustrophobia. Thus, a complete 1 year follow-up including clinical and MRI investigation was performed in 10 patients (90%).

Intraoperative findings. The arthroscopic evaluation of the eleven patients revealed nine type III (82%) and two type IV (18%) SSC lesions. Thus, all patients showed at least complete lesions of the tendon's superior two-thirds with some retraction from intermediate up to the level of the glenoid rim. In three cases, minor PASTA lesions were seen. However, these lesions did not require any surgical treatment due to their small size involving only the innermost part of the deep tendon layer.

The LHBT was completely dislocated in three patients. Subluxation of the LHBT with the tendon riding on the anterior aspect of the bicipital groove was seen in five cases (45%), whereas two biceps tendons (18%) presented with a pronounced anterior instability due to a lesion to the anterior pulley. A concomitant SLAP lesion was diagnosed in only one patient (9%). In two patients a partial tear $< 50\%$ of the biceps tendon was evident (18%). Only one LHBT was considered as normal.

Follow-up. One year postoperatively, nine patients (82%) were back at their previous work. One patient was already retired at the time of the injury and one patient was already disabled due to a cervical spine injury. Six patients reported a full return to their sports activities whereas one patient had to reduce his sports due to some persistent shoulder problems. The remaining four patients had never participated in any sports activities, not even before the injury.

Clinical outcome. The results of the clinical examination are shown in table 1. Preoperatively, all patients were able to perform the belly-press test. However, muscular strength was reduced in all patients compared to the contralateral side. ~~However, the test was considered positive in all patients.~~ Seven patients (64%) presented with a positive lift-off test, in one case a lag sign was evident. Four patients (36%) failed to demonstrate a correct lift-off test preoperatively due to limited internal rotation or pain exacerbation. In contrast, at follow-up all patients correctly performed both tests and a significant improvement of these specific SSC tests regarding strength was found. Compared to the uninjured contralateral side, the mean loss of internal rotation was measured 1.5 ± 2.0 points (0-6) in the CMS subscore for internal rotation ($p < 0.05$). External rotation in 0° abduction was $46 \pm 19^\circ$ (20-70 $^\circ$) on the operated side compared to $57 \pm 18^\circ$ (30-85 $^\circ$) on the contralateral uninjured side ($p < 0.05$). Strength of external rotation was similar to the uninjured side. The CMS and all its subscores, such as activities of daily living, pain, range of motion and strength demonstrated a marked improvement at follow-up when compared to the preoperative situation. Nine patients rated their outcome excellent (82%), one patient had a good result and one patient was satisfied.

MRI follow-up. Complete structural integrity of the SSC tendon was seen in all investigated patients ($n=10$). Neither a partial nor a complete re-rupture of the reconstruction was observed. Alterations of the muscular mass and the course of the fatty infiltration of the SSC are shown in table 2. The CSA of the SSP and ISP significantly increased although there was neither a significant lesion found arthroscopically nor were these muscles/tendons involved in any surgery.

Interestingly, there was no change in fatty infiltration in these muscles. No failure of the LHBT tenodesis was observed and all the humeral heads were anatomically centered in the glenoid.

Discussion

Arthroscopic repair of SSC lesion Lafosse III and IV is associated with a good clinical outcome including high patient satisfaction. MRI follow-up at one year demonstrated complete healing of the reconstructed tendon with no re-rupture.

Burkhart⁵ published the first article describing arthroscopic SSC tendon repair in 2002. Still, the evidence for arthroscopic repair of isolated SSC tears is low. A recent systematic review comparing open and arthroscopic surgical repair of isolated SSC lesions failed to demonstrate a clear advantage for either method^{4,6,18}. However, one must realize that only level IV studies were available for this evaluation. Good pain relief and excellent function may be achieved by open and arthroscopic surgery. Edward et al.³ published the largest series so far describing open repair of isolated SSC tears, either of traumatic origin or due degeneration. Of these, 23 were limited to the superior one-third of the tendon leaving 64 patients with higher-grade lesions.

Furthermore, there are only a few studies available with a radiological follow-up by either CT arthrography or MRI^{1, 7-10,11}. In contrast to other studies, we focused on isolated traumatic SSC lesion with at least moderate tendon retraction.

Clinical outcome. In general, with either open or arthroscopic repair of the SSC lesion good clinical results can be achieved. In accordance with the literature, we observed a significant improvement of the CMS and all its subcategories. CMS achieved after arthroscopic repair of isolated SSC lesion ranges from 74 to 85^{9,10,11}. In our series, CMS after one year was 90. One could hypothesize that this result was achieved due to short interval between trauma and surgery. This view is supported by the observation that a delay in open SSC repair resulted in poorer

clinical outcome^{3,4}. In accordance with the CMS, the specific SSC tests such as the modified belly-press- and the lift-off test also improve significantly after surgery^{1, 7- ~~10~~11}.

LHBT pathologies. Beside the SSC tendon repair, we also performed an anchor tenodesis in all patients no matter what the underlying pathology was. At follow-up, all tenodeses were intact. In a series with 40 patients with isolated or combined rotator cuff tendon lesions a concomitant pathology of the LHBT was found in 63%⁶. In the largest series investigating open repair of isolated SSC tears, Edwards et al.³ could clearly demonstrate that the performance of either LHBT tenodesis or tenotomy had a beneficial effect on the CMS and the subjective outcome regardless of the preoperative condition of the LHBT. Thus, the authors suggested to perform routine LHBT tenodesis or tenotomy at the time of the SSC repair.

Structural integrity of the SSC tendon repair. In a human cadaver study Wellmann et al.^{~~17~~-19} compared single-row repair with a double row repair using a “suture bridge technique” as well. The double-row technique restored 48% of the ultimate load of an intact tendon, whereas a single-row repair failed significantly earlier at 34%. At one-year follow-up, all SSC tendon repairs of our study group were intact as no re-rupture was evident on MRI one year postoperatively. We routinely performed a “pseudo double-row” repair for Lafosse III tears. A double-row repair with suture bridges was used for more retracted IV lesions. According to the current literature re-rupture rates range from 5-14%^{1, 7, ~~8-9~~, ~~10~~11}. In all these studies either CT arthrography or MRI studied structural integrity at follow-up 20-~~36-57~~ months postoperatively. In the largest published study so far, Toussaint et al.^{~~9~~-10} could evaluate 129 patients with either isolated lesions or in combination with SSP tears with a radiological follow-up of at least 6 months. Re-rupture rate in this large series was 8%.

Alterations of the SSC muscle. Although MR evaluation demonstrated a significant decrease in fatty SSC muscle infiltration and an increase of muscular mass at follow-up compared to the preoperative images, the interpretation of these findings may be controversial. Our observations

are in contrast to the results of some other studies found in the recent literature ^{9-10,11} which all reported a progression of fatty infiltration despite successful surgery. Interestingly, no correlation with clinical outcome was found. Only Bartl et al. ⁷ and Lafosse ¹ did not observe a progression of fatty infiltration. However, interval between trauma/onset of symptoms and surgery ranged from 5.8 months ⁷ up to 35.7 months ^{10,11} in these studies. One could hypothesize that in our study, the SSC muscle improved in quantity (mass) and quality (less fatty infiltration) due to the fact that the mean interval from trauma to surgery was only 3.7 months and thus considerably shorter than in most other studies ^{1, 5, 7, 8, 10,11}. Interestingly, fatty degeneration was already evident on the preoperative images of our study group despite the short interval between trauma and diagnostic work-up. One could hypothesize that fatty degeneration may occur earlier and develop quicker in traumatic lesions than in degenerative SSC tears. However, to our knowledge, there is no scientific data available to support this clinical observation.

Our findings may indicate that muscular changes of the SSC are reversible if the tendon is reattached shortly after trauma. However, the significance of fatty degeneration and muscle wasting is still unclear. In their multicentre study, Toussaint et al. ⁹⁻¹⁰ reported no adverse clinical effects despite marked muscle alterations. Furthermore, the comparison of the muscle mass and fatty degeneration pre- and postoperatively may be hazardous since the noted differences could also be attributed to a pure volumetric distortion once a retracted muscle is reattached. Moreover, preoperative MRI was contrast-enhanced, the follow-up MRI was not. This fact may have impaired the accuracy of our measurements. This view is supported by the fact that we also observed an increase of the CSA for the SSP and ISP although they were neither injured nor involved in any surgery. However, fatty infiltration was decreased in the SSC at follow-up whereas it remained unchanged in the other two studied muscles. In a recent study Jo et al. ^{18,20} compared preoperative MRI investigations to MRI studies obtained 3 days following surgery. In a first study, a significant increase of the CSA of the SSC was demonstrated despite no surgical cuff repair was performed in these patients. No conclusive reason for this finding was given by the authors. Thus, the CSA of the SSC was excluded from further evaluations. However, no difference was seen

regarding fatty infiltration of the SSC and no difference was found in the SSP and ISP regarding both parameters. In a second study, patients with arthroscopic rotator cuff repair were evaluated. Interestingly, rotator cuff repair significantly increased the CSA of the SSP by as much as 45% for massive tears. The decrease of the fatty infiltration was significant as well, but only for the SSP and ISP whereas no significant change was found for the SSC. To overcome this potential bias, the authors suggested that images should also be obtained immediately after the surgical procedure and compared with long-term follow-up for a true assessment of muscular alterations rather than being compared with preoperative images.

Limitations of the current study

With ten patients available for a complete follow-up and one patient with a clinical follow-up only, the sample size of this case series is small and the study design lacks a comparative control group. We studied only complete isolated SSC tears (Lafosse III and IV) which are usually considerably retracted. Since we did not include minor lesions such as Lafosse I and II, our results are not falsely improved by the inclusion of clinically less significant tears. Furthermore, anterosuperior tears and mass lesions of the rotator cuff were not included.

In the largest study published so far a total of 208 SSC tears, either isolated or associated with a limited anterosuperior lesion, were analyzed^{9,10}. Of these, only 35 patients with isolated or “very predominant” subscapularis lesions were available for follow-up. Heikenfeld et al.⁸ published a case series with 20 patients. However, they also included 10 Lafosse II lesions. Other authors published case series with similar sample sizes ranging from 7 to 17 patients when only tears equivalent to Lafosse III and IV lesions were counted^{1, 7, 11+9}. In a recently published study, 46 patients with arthroscopic repair of only large SSC lesions (Lafosse III-IV) were investigated⁹. However, only 6 SSC tears were isolated, either traumatic or of degenerative nature.

Furthermore, a thorough comparison between different studies may be compromised to different inclusion criteria, the variety of different classification systems to grade SSC lesions, the measured outcome parameters, the duration of follow-up and the low evidence (Level IV) of the available literature¹⁶¹⁸.

Conclusions

Arthroscopic repair of higher grade isolated SSC lesions provides reliable tendon healing accompanied by excellent functional results one year after surgery. ~~In our series we observed significant decrease in fatty SSC muscle infiltration and an increase of muscular mass.~~

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Legends

Figure 1. Lesion of the subscapularis tendon (SSC) Type Lafosse III (right shoulder). **A.** Lesion before repair (view through the posterior A portal). **B.** Subscapularis tendon with applied traction suture for better mobilization and reduction (view through the anterolateral D portal). SGHL, superior gelnohumeral ligament. HH, humeral head.

Figure 2. Visualisation and identification of the axillary nerve (AN) during mobilization of the subscapularis muscle (right shoulder, view through the anterolateral D portal).

Figure 3. Left shoulder, view through the dorsal A portal. Situation with applied anchors and sutures before final reduction and knot tying. The reconstruction is performed using U-stitches and a single lasso-loop stitch (*)-(black arrow) at the upper border of the subscapularis tendon (SSC). For better orientation violet and blue different colored strands are used. The traction suture (green-strand, **)-white arrow) is removed before completion of the reconstruction.

Figure 4. A. Axial preoperative MR arthrography slice demonstrating lesion of the subscapularis tendon (*)-(SSC, white arrow) with anterior dislocation of the long head of biceps tendon (**)- (black arrow). **B.** Corresponding follow-up MR slice showing full structural integrity of the subscapularis repair (white arrow) -(#) and tenodesis of the long head of biceps tendon (\$)-(black arrow).

Figure 5. A. Preoperative axial MR slice with fatty infiltration of the subscapularis muscle (white arrow) -(*). **B.** The corresponding MR slice at one year follow-up shows a marked reduction of the fatty infiltration. SSC, subscapularis muscle; SSP, supraspinatus muscle; ISP, infraspinatus muscle. (**).

Table I Clinical Results

	Preoperative	1 year follow-up	P value
Modified belly-press test (strength, max 5 points)	2.9±0.3 (2-3)	4.8±0.6 (3-5)	< 0.05
Modified lift-off test (strength, max 5 points)	2.9±0.4 (2-3)	4.8±0.6 (3-5)	< 0.05
Internal rotation (strength, max 5 points)	3.1±1.6 (0-6)	7.8±2.6 (2-10)	< 0.05
CMS total (max 100 points)	43.5±21.3 (16-80)	89.3±15.0 (51-100)	< 0.001
CMS ADL (max. 20 points)	8.2±4.8 (2-18)	18.4±3.1 (10-20)	< 0.05
CMS Pain (max 15 points)	4.6±4.2 (0-10)	13.2±3.8 (5-15)	< 0.05
CMS ROM (max 40 points)	22.7±7.2 (12-32)	36.0±5.4 (22-40)	< 0.05
CMS Strength (max 25 points)	8.0±7.0 (0-20)	21.7±5.3 (10-25)	< 0.05

430

431 CMS, Constant-Murley score; ADL, activities of daily living; ROM, range of motion; Data given as

432 mean with standard deviation, the range is provided in brackets.

433

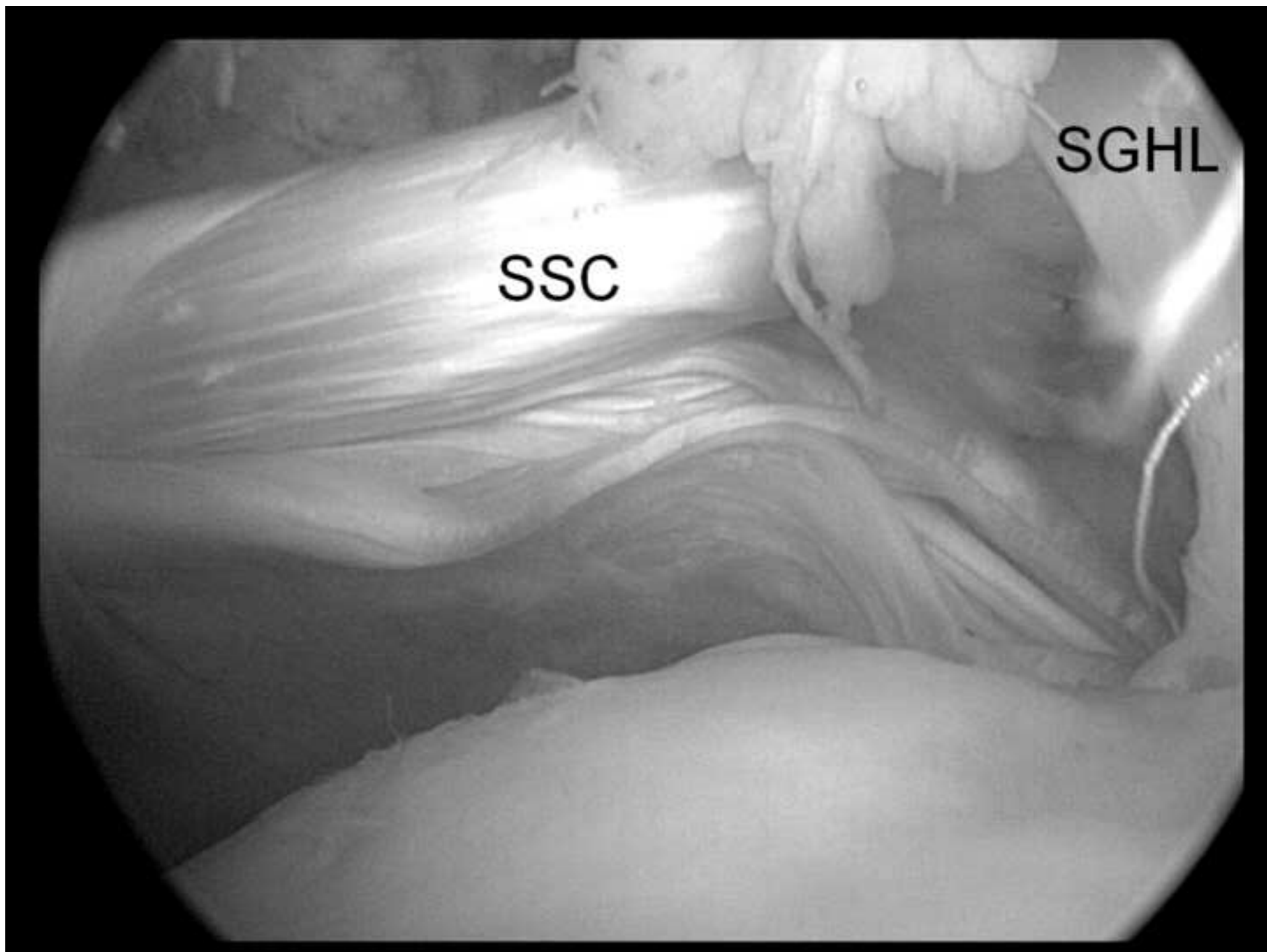
Table II MRI evaluation: Fatty infiltration and CSA of the rotator cuff muscles

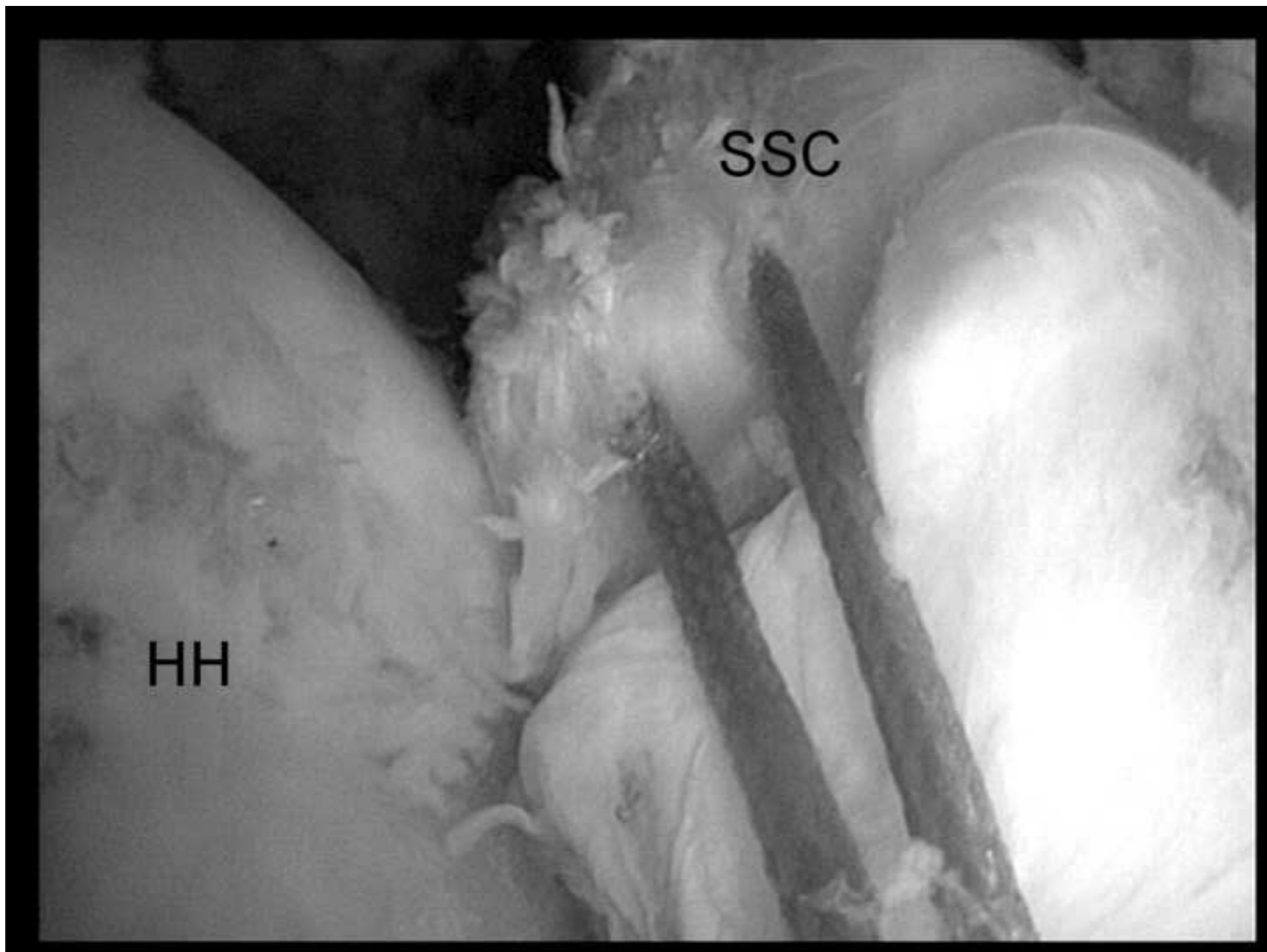
	Preoperative	1 year follow-up	P value
Fatty infiltration of SSC	1.7 (0-3)	0.7 (0-2)	< 0.05
CSA SSC (mm ²)	1491 (900-2120)	2158 (1370-3080)	< 0.001
Fatty infiltration of SSP	0.4 (0-2)	0.2 (0-2)	n.s.
CSA SSP (mm ²)	682 (350-940)	865 (480-1060)	< 0.001
Fatty infiltration of ISP	0.3 (0-2)	0.3 (0-2)	n.s.
CSA ISP (mm ²)	1441 (990-1830)	1716 (1020-2300)	< 0.001

434

435 | Fatty infiltration of the SSC was graded according to Goutallier ¹³¹⁴, modified by Fuchs ¹⁴¹⁵. The
436 | cross-sectional area of the SSC was measured according to the method proposed by Zanetti et al.
437 | ¹⁵¹⁶. SSC, subscapularis muscle; SSP, supraspinatus muscle; ISP, infraspinatus muscle; CSA,
438 | cross sectional area; data given as mean with range in brackets

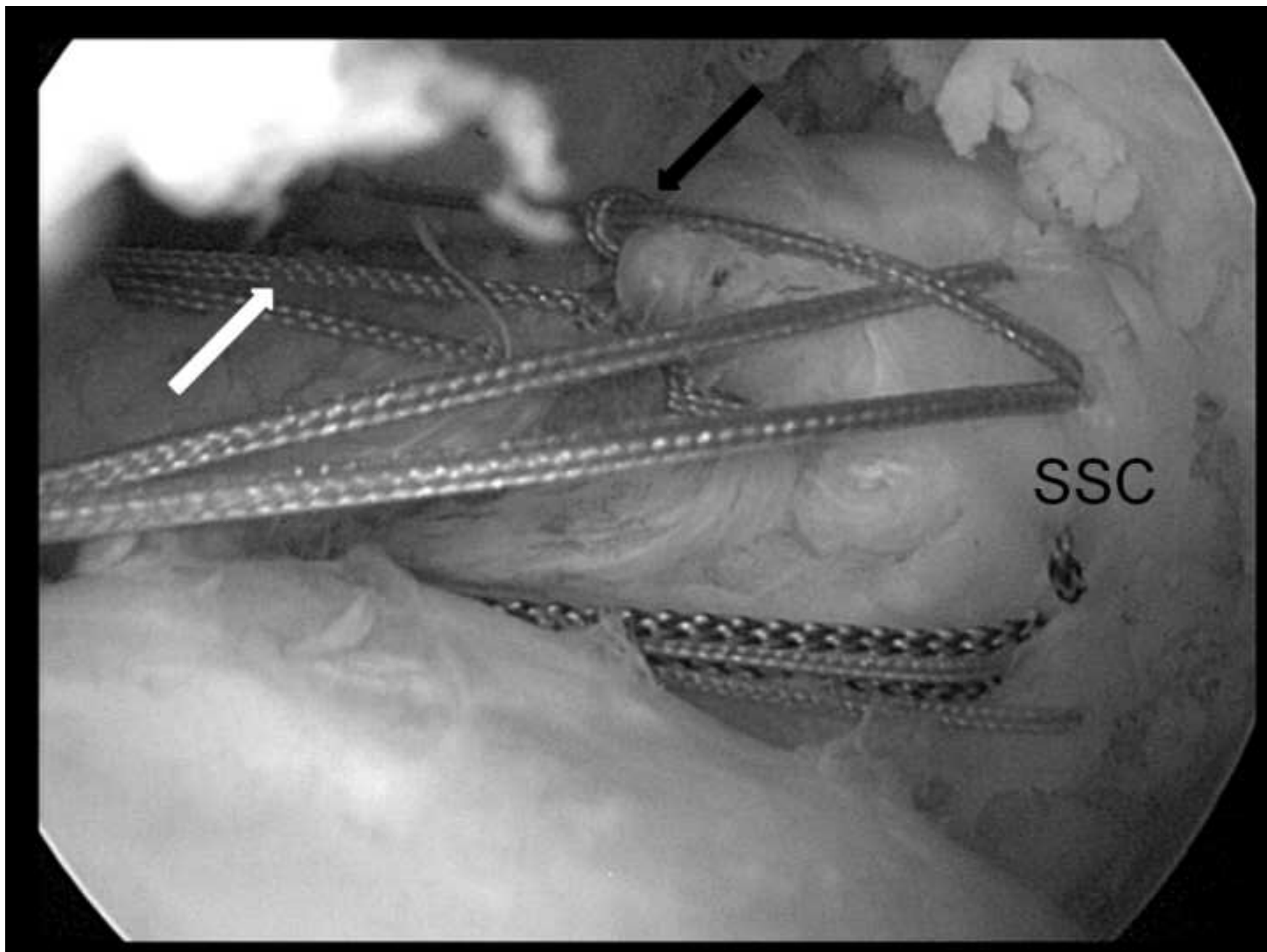
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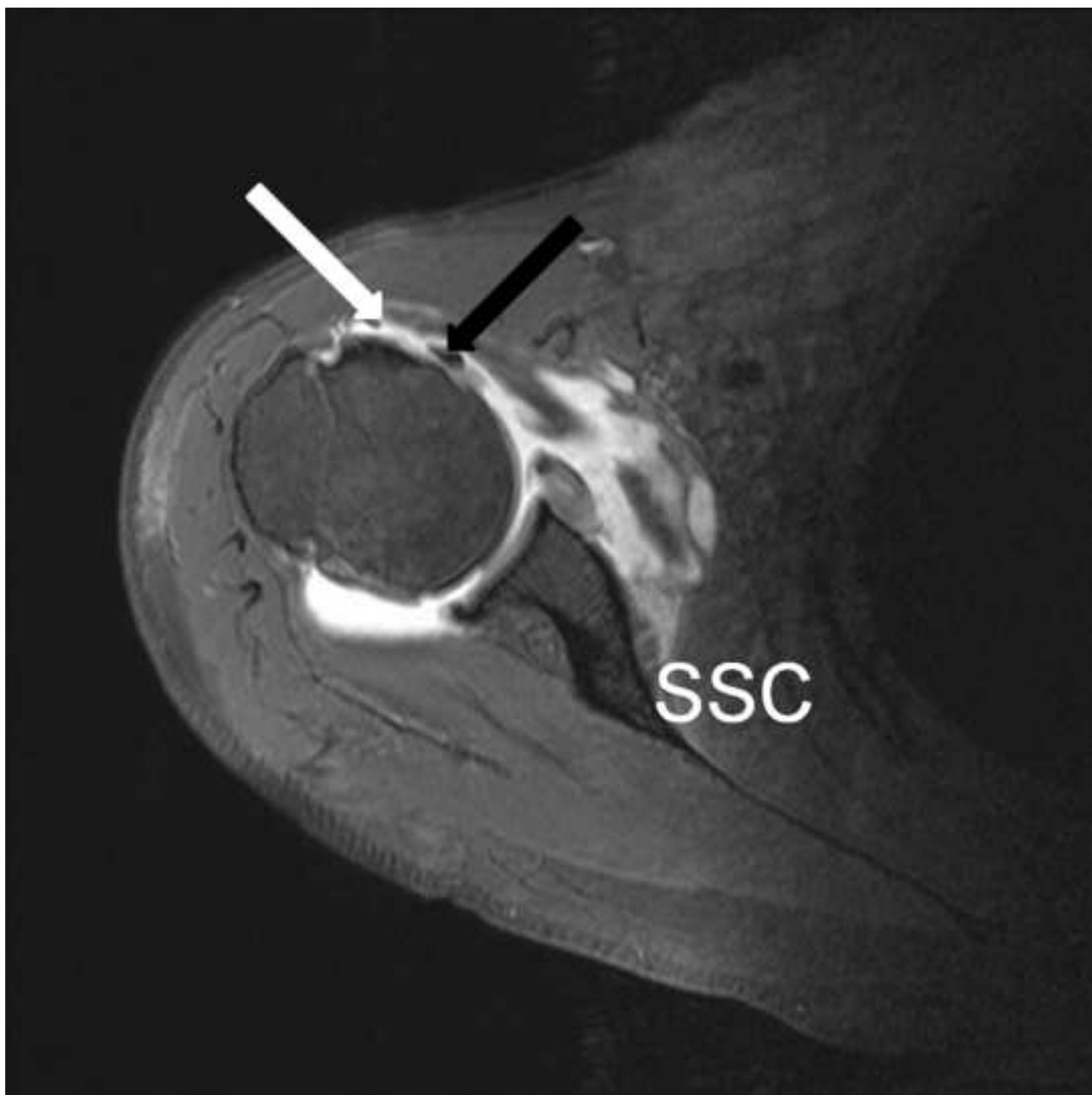


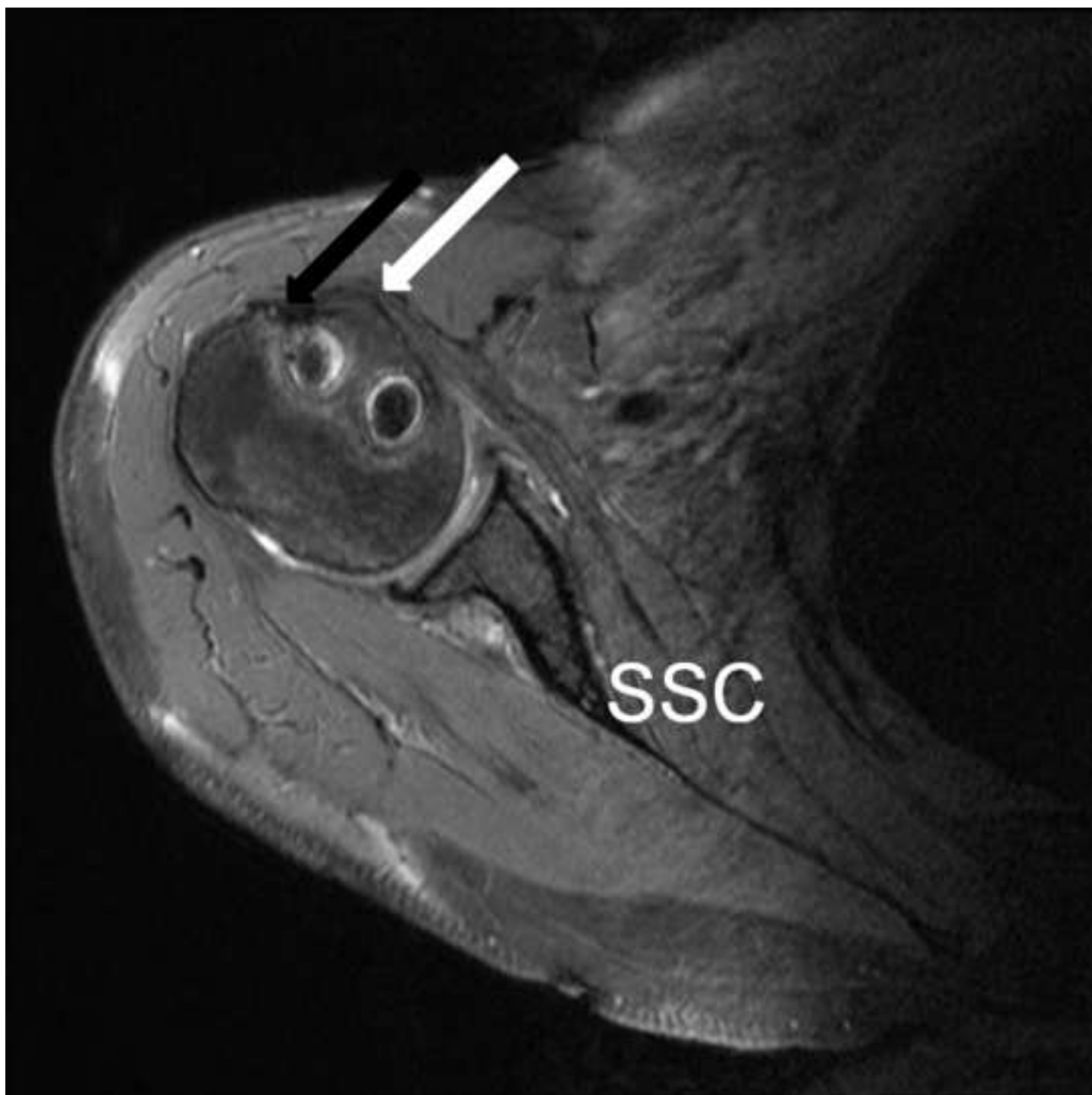


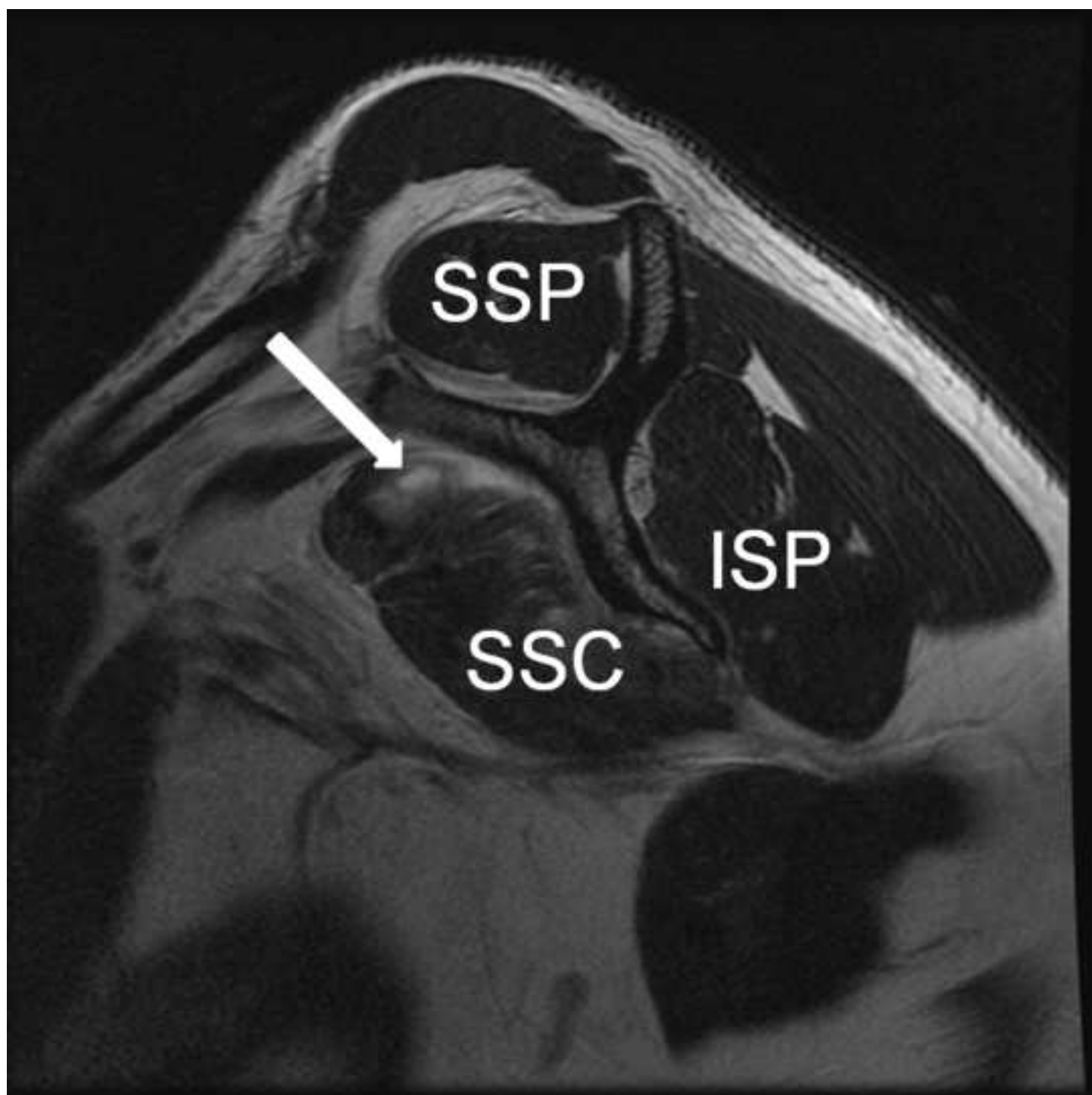


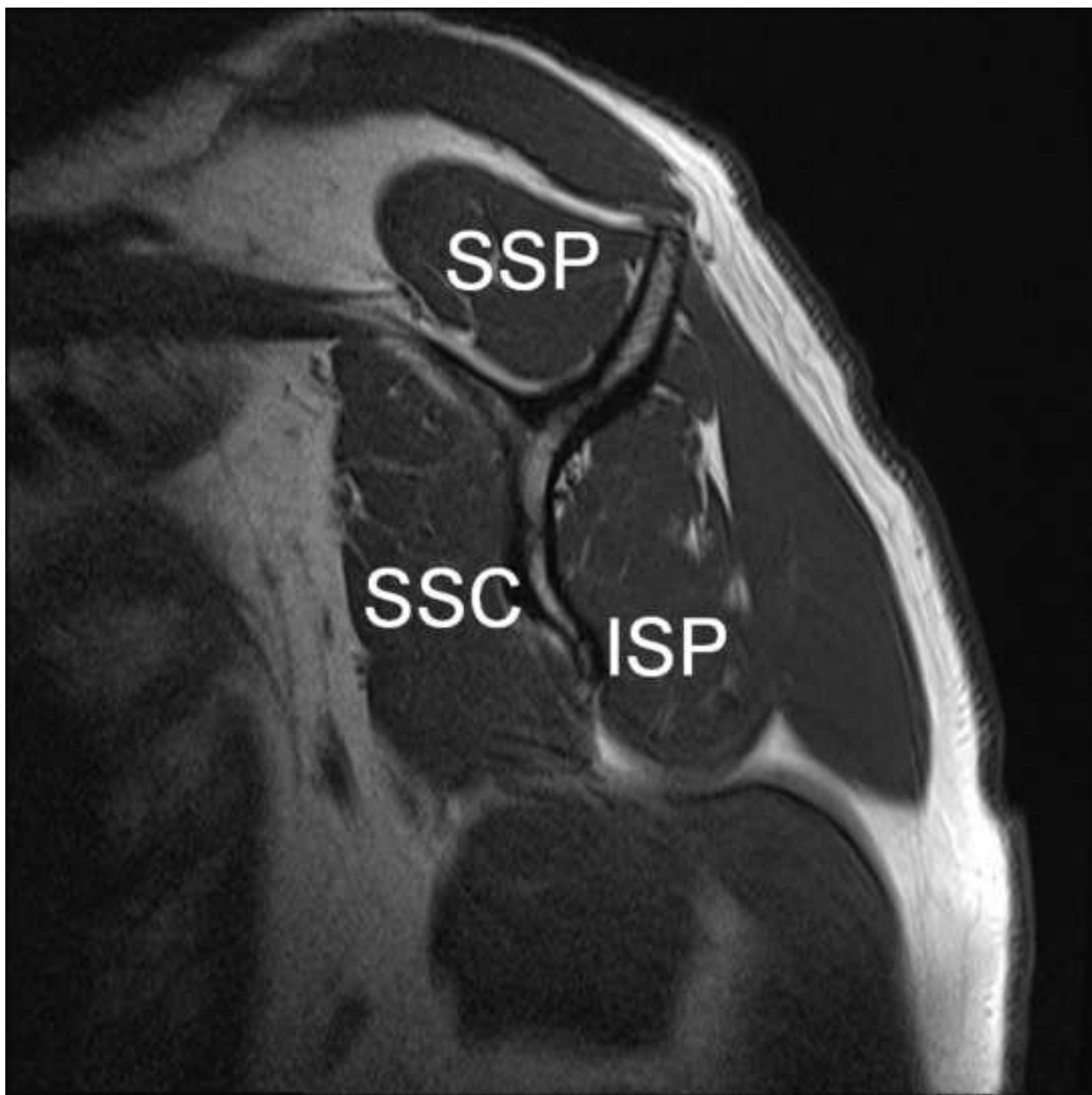
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To
Mr. G.G. Poehling, MD
Editor-in-Chief Intensive Arthroscopy
CompRehab Plaza
131 Miller Street
Winston-Salem, NC 27103
USA

January 16, 2014

Dear Sir

Please find enclosed our revised version of the manuscript

“Arthroscopic repair of traumatic isolated subscapularis tendon lesions Lafosse III-IV: a prospective MRI-controlled case series with one year follow-up”

Ref.: ARTH-13-784

by Patrick Grueninger et al.

which we like to resubmit for publication in *Arthroscopy: The Journal of Arthroscopic and Related Surgery*

We have appreciated the fair and constructive criticisms of the reviewers. In the following, please find our point-by-point reply to the reviewer's comments. Page and line numbers of changes made refer to the revised version of the manuscript to facilitate reading.

Associate Editor's Comments:

Authors, you should be commended on the quality of your scientific work. Your study has strict inclusion criteria with a well defined patient group. Your outcome measures are chosen appropriately and you have included imaging follow-up. However, the main limitation to your study is that the information is not new, and has been presented in the literature before, most recently by Lafosse in JBJS. With the small patient group you have reported on, there is no new information that is presented for the reader. The impact to clinical practice is limited.

Response: Thank you very much for your comment. We fully agree with your points and we are fully aware of our work's limitations. To overcome some of them, we have applied very strict inclusion criteria to enhance the scientific information provided by our study. In contrast to the study by Lafosse et al. (1), published in 2007, we have only included traumatic SSC lesions Lafosse III-IV. Non-traumatic lesions and low-grade lesions (I-II) were excluded.

The most recent study was published in *Arthroscopy* by L. Lafosse and his group in September 2013 (2). We were not aware of this publication when our manuscript was submitted. Thus, we have included this reference in our revised version of the manuscript. Accordingly, the following sentences were inserted into the *Discussion* section (see lines 349-351 in the revised manuscript). Furthermore, the references were adapted, accordingly.

1. Lafosse L, Jost B, Reiland Y, Audebert S, Toussaint B, Gobezie R. Structural integrity and clinical outcomes after arthroscopic repair of isolated subscapularis tears. *J Bone Joint Surg Am* 2007; 89: 1184-1193.

2. Lanz U, Fullick R, Bongiorno V, Saintmard B, Campens C, Lafosse L. Arthroscopic large subscapularis tendon tears: 2- to 4-year clinical and radiographic outcomes. *Arthroscopy* 2103; 29(9): 1471-1478.

Reviewer #1:

Comment 1: The introduction is well written and concisely states the purpose of the study. The authors point out the paucity of information on this subject in the literature. Line 36 "rare" should be replaced by "less common".

Response: The term "rare" is replaced by "less common" (see line 40 in the revised version of the manuscript).

Comment 2: The Methods are well described and do answer the central question. One criticism is the inclusion of determining reversal of fatty infiltration. If these are indeed acute traumatic events then there should be little if any fatty infiltration in the mean average of 3.7 months from injury to surgery. This should be eliminated.

Response: Thank you for this important comment. All our study patients sustained a trauma and there is no history of previous shoulder pain or impaired function. However, we do not know why some MR studies demonstrated fatty infiltration of the SSC so early after trauma. One could speculate that fatty degeneration may develop earlier in patients with large traumatic lesions than in cases with chronic degeneration. Furthermore, this fatty infiltration may be reversed when tendon repair is performed early enough. This view is derived from our clinical experience. However, this is pure speculation and to our knowledge there is no scientific data available neither to support nor to contradict this hypothesis. Nevertheless, our MR images were thoroughly analyzed and the occurrence of fatty infiltration was already evident on the preoperative MR images. Furthermore, statistically significant decrease of fatty degeneration was found at follow-up indicating that this process may be reversible. Thus, we do not think this data/information should be eliminated. However, we have included this important aspect in the *Discussion* section (see lines 305-309 in the revised manuscript).

Comment 3: All references to specific anchors and manufacturers should be replaced by "threaded suture anchors" since this paper is about success of surgery not testing specific anchors.

Response: These terms were corrected accordingly throughout the revised version of the manuscript (see line 113 and 117).

Comment 4: Muscle strength was grade from 0 to 5 according to the classification of neurological assessment. This is too subjective for inclusion in this study. Were the pre and post-op examinations performed by the same person?

Response: The clinical examinations were all performed by the first author (see line 139 in the revised manuscript) with exactly the same technique and muscular strength was compared with the uninjured contralateral shoulder. We absolutely agree with the reviewer's view that the classification of neurological assessment has its limitations regarding an accurate assessment of muscle strength. It would have been more accurate and objective to use a spring gauge. However, this grading system has also been used by other investigators for similar purposes such as Lafosse (2). Others (3) only graded between negative, asymmetric and positive. Thus, we considered the classification of neurological assessment appropriate when the current study was designed. We will certainly consider another measuring method for future research projects.

2. Lafosse L, Jost B, Reiland Y, Audebert S, Toussaint B, Gobezie R. Structural integrity and clinical outcomes after arthroscopic repair of isolated subscapularis tears. *J Bone Joint Surg Am* 2007; 89: 1184-1193.

3. Nové-Josserand L, Hardy M-B, Ogassawara RLN, Carrillon Y, Godenèche A. Clinical and structural results of arthroscopic repair of isolated subscapularis tear. *J Bone Joint Surg Am* 2012; 94: e125 (1-7)

Comment 5: Why were "routine acromioplasties" performed? Were coracoplasties performed?

Response: An acromioplasty was performed in all patients as a matter of routine. All our patients undergoing therapeutic shoulder arthroscopy receive subacromial debridement and an acromioplasty at the end of the procedure. However, in asymptomatic patients without a spur, we perform a very limited instead of a formal acromioplasty as it was the case in this study group (see lines 119-120 in the revised version of the manuscript). We did not perform any coracoplasties in this study group as we did not see any coracoid spurs or signs of impingement. None of our patients presented with a type V SSC tears (Lafosse classification) which would be associated with an eccentric head causing coracoid impingement. This aspect is now included in the revised version of the manuscript (see lines 118-119).

Comment 6: Was the radiologist MSK fellowship trained or a generalist? Did the same radiologist evaluate the pre and post-op studies?

Response: All pre- and postoperative MR studies were evaluated by the same radiologist who indeed is experienced and properly trained in musculoskeletal imaging techniques. He has repeatedly attended specialty courses for MSK radiology. This information was added in the revised version of the manuscript (see lines 174-175).

Comment 7: Pre-op MRIs were performed with intraarticular gadolinium whereas the post-op studies were performed without contrast. The authors need to address this discrepancy and how it skews to results.

Response: Preoperatively, all patients underwent MRA as the standard radiologic investigation technique for patients with suspected injury to the rotator cuff or SLAP lesions at our institution.

Our patients did all well accept arthrography to investigate their injured shoulder for a proper diagnosis and the planning of the following therapeutic steps. Since the number of our study group is quite small, we strongly depended on a complete or at least near complete follow-up of our patients. Acceptance of an invasive technique for pure scientific reasons without direct benefit to the patient may be low and may also be discussed controversially by ethical aspects. Furthermore, the superiority of MRA for evaluating structural integrity of SSC repair is not proven (4). Thus, only MR imaging was performed at follow-up. All SSC lesions were arthroscopically graded, not by MRA. Fatty infiltration and the cross-sectional area of the SSC were evaluated by MRA preoperatively and MRI at follow-up, respectively. However, the application of intra-articular gadolinium does not change the appearance of the muscles such as the SSC on the images and direct comparison may be eligible. This information including the reference (4) is now given in the revised version of the manuscript (see lines 168-173).

4. Duc SR, Mengiardi CWA, Jost B, Hodler J, Zanetti M. Diagnostic performance of MR arthrography after rotator cuff repair. *AJR* 2006; 186: 237-241.

Comment 8: The authors need to explain their statement "However, the test [belly press] was considered positive in all patients" and how this correlates to excellent results.

Response: All patients were able to perform the belly-press test. However, muscular strength was impaired in all patients as well. This is also shown in table 1. The sentence "However, the test was considered positive in all patients" is not correct. We apologize for this misleading expression. Our patients could perform the belly-press but not with the same strength as on the contralateral side. This mix-up is now corrected in the revised version of the manuscript (see lines 119-121).

Comment 9: The first part of the conclusion statement is supported by the evidence. The second part is clearly not supported and should be eliminated.

Response: We agree with the reviewer's opinion. Thus, the second part of the Conclusion was eliminated in the *Conclusion* section and the *Abstract* as well (see lines 32 and 360).

Comment 10: I concur this is a Therapeutic Level IV study.

Response: Yes. This information is given in the abstract (see line 32). Does it need to be mentioned anywhere else in the paper?

Comment 11: The title is adequate although not completely descriptive.

Response: The title was left unchanged.

Comment 12: The operative photos are identified in the legends but would be more readable if the photos themselves were labeled. Color photos are not necessary for publication. The MRI images should also be labeled.

Response: All color photos have been converted to black and white images in the revised version of the manuscript. Furthermore, all images were labelled and the legends adapted

accordingly (see lines 424-445 in the revised version of the manuscript and all revised figures (Fig 1A – 5B)).

Reviewer #2:

Comment 1: Line 38: Delete over the last years.

Response: This term is eliminated in the revised version of the manuscript (see line 42).

Comment 2: Line 45: outcomes.

Response: This typo is corrected (see line 49).

Comment 3: Line 60: Was this case series done retrospectively or prospectively?

Response: It is a prospective case series. This is now mentioned in the *Methods* section of the manuscript (see line 69).

Comment 4: Line 170: Claustrophobia?

Response: The reviewer is right, of course. We apologize for this embarrassing mix-up: “agoraphobia” was changed to “claustrophobia” (see line 195 in the revised version of the manuscript).

Comment 5: The title is too long in my opinion.

Response: We have considered a shorter title. However, we feel, that the original title quite accurately describes what the study is all about. With a shorter title some of this information would be lost and a mix-up with other studies such as the one mentioned below (1) would be more likely.

1. Lanz U, Fullick R, Bongiorno V, Saintmard B, Campens C, Lafosse L. Arthroscopic large subscapularis tendon tears: 2- to 4-year clinical and radiographic outcomes. *Arthroscopy* 2103; 29(9): 1471-1478.

Overall, the authors feel that the quality of the manuscript has markedly improved due to the changes made according to the suggestions of the reviewers, and we would be happy if it now meets the criteria for publication in *Arthroscopy: The Journal of Arthroscopic and Related Surgery*.

Sincerely yours

Patrick Grueninger
Christoph Meier

Arthroscopy: The Journal of Arthroscopic and Related Surgery

— Instructions —

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Each author of the manuscript must separately complete and save this form using his or her name in the file name. Each author's completed form must then be uploaded with the manuscript.

The purpose of this form is to provide readers of your manuscript with information about your other interests that could influence how they receive and understand your work. The form is in four parts:

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Enter your full name and provide the manuscript title.

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This section asks for information about the work that you have submitted for publication. The time frame for this reporting is that of the work itself, from the initial conception and planning to the present. The requested information is about resources that you received, either directly or indirectly (via your institution), to enable you to complete the work. Checking "No" means that you did the work without receiving any financial support from any third party — that is, the work was supported by funds from the same institution that pays your salary and that institution did not receive third-party funds with which to pay you. If you or your institution received funds from a third party to support the work, such as a government granting agency, charitable foundation, or commercial sponsor, check "Yes." Then complete the provide the information requested.

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This section asks about your financial relationships with entities in the biomedical arena that could be perceived to influence, or that give the appearance of potentially influencing, what you wrote in the submitted work. You should disclose interactions with ANY entity that could be considered broadly relevant to the work.

Report all sources of revenue paid (or promised to be paid) directly to you or your institution on your behalf over the 36 months prior to submission of the work. This should include all monies from sources with relevance to the submitted work, not just monies from the entity that sponsored the research. Please note that your interactions with the work's sponsor that are outside the submitted work should also be listed here. If there is any question, it is usually better to disclose a relationship than not to do so.

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-

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3. Employment

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4. Expert testimony

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5. Grants/grants pending

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6. Payment for lectures including service on speakers bureaus

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7. Payment for manuscript preparation

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8. Patents (planned, pending or issued)

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9. Royalties

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10. Payment for development of educational presentations

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11. Stock/stock options

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** For example, if you report a consultancy above there is no need to report travel related to that consultancy on this line.

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Are there other relationships or activities that readers could perceive to have influenced, or that give the appearance of potentially influencing, what you wrote in the submitted work?

 X No other relationships/conditions/circumstances that present a potential conflict of interest

___ Yes, the following relationships/conditions/circumstances are present (explain below):

>>>>>>>>>>><<<<<<<<<<<<

The International Committee of Medical Journal Editors

The ICMJE Disclosure of Potential Conflicts of Interest Form was adopted by *Arthroscopy: The Journal of Arthroscopic and Related Surgery* along with 17 other leading orthopaedic journals at the 2011 annual meeting of the American Academy of Orthopaedic Surgeons.

ICMJE Form for Disclosure of Potential Conflicts of Interest

Section 1. Identifying Information

1. Given Name Joerg 2. Surname Schneider
3. Are you the corresponding author? Yes ___ No ☒ X
4. Effective Date 5/11/13
5. Manuscript Title Arthroscopic repair of traumatic isolated subscapularis tendon lesions Lafosse III-IV: a prospective MRI-controlled case series with one year follow-up
-

Section 2. The Work Under Consideration for Publication

Did you or your institution at any time receive payment or services from a third party for any aspect of the submitted work (including but not limited to grants, data monitoring board, study design, manuscript preparation, statistical analysis, etc...)?

Complete each item by typing an X in answer yes or not and completing the information requested if an answer is Yes. If you have more than one relationship, add lines.

1. Grant

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13. Other (err on the side of full disclosure)

☒ No ___ Yes, money paid to you ___ Yes, money paid to institution* Name of entity___ Comments___

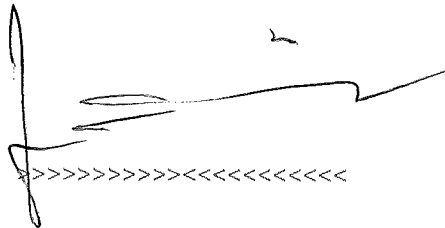
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Section 4. Other relationships

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☒ No other relationships/conditions/circumstances that present a potential conflict of interest

☐ Yes, the following relationships/conditions/circumstances are present (explain below):

A handwritten signature in black ink, consisting of a vertical line on the left and a horizontal line extending to the right, with a small flourish at the end.

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Section 1. Identifying Information

1. Given Name Andreas_ 2. Surname Platz
 3. Are you the corresponding author? Yes No ☒ X
 4. Effective Date 04 November 2013
 5. Manuscript Title Arthroscopic repair of traumatic isolated subscapularis tendon lesions Lafosse III-IV: a prospective MRI-controlled case series with one year follow-up
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Enter your full name and provide the manuscript title.

Section 2. The work under consideration for publication

This section asks for information about the work that you have submitted for publication. The time frame for this reporting is that of the work itself, from the initial conception and planning to the present. The requested information is about resources that you received, either directly or indirectly (via your institution), to enable you to complete the work. Checking "No" means that you did the work without receiving any financial support from any third party — that is, the work was supported by funds from the same institution that pays your salary and that institution did not receive third-party funds with which to pay you. If you or your institution received funds from a third party to support the work, such as a government granting agency, charitable foundation, or commercial sponsor, check "Yes." Then complete the provide the information requested.

Section 3. Relevant financial activities outside the submitted work

This section asks about your financial relationships with entities in the biomedical arena that could be perceived to influence, or that give the appearance of potentially influencing, what you wrote in the submitted work. You should disclose interactions with ANY entity that could be considered broadly relevant to the work.

Report all sources of revenue paid (or promised to be paid) directly to you or your institution on your behalf over the 36 months prior to submission of the work. This should include all monies from sources with relevance to the submitted work, not just monies from the entity that sponsored the research. Please note that your interactions with the work's sponsor that are outside the submitted work should also be listed here. If there is any question, it is usually better to disclose a relationship than not to do so.

For grants you have received for work outside the submitted work, you should disclose support ONLY from entities that could be perceived to be affected financially by the published work, such as entities or foundations supported by entities that could be perceived to have a financial stake in the outcome. Public funding sources, such as government agencies, charitable foundations, or academic institutions, need not be disclosed. For example, if a government agency sponsored a study in which you have been involved and drugs were provided by a pharmaceutical company, you need only list the pharmaceutical company.

Section 4. Other relationships

Use this section to report other relationships or activities that readers could perceive to have influenced, or that give the appearance of potentially influencing, what you wrote in the submitted work.

ICMJE Form for Disclosure of Potential Conflicts of Interest

Section 1. Identifying Information

1. Given Name Patrick_ 2. Surname Grueninger
3. Are you the corresponding author? Yes ☒ No
4. Effective Date 04 November 2013
5. Manuscript Title Arthroscopic repair of traumatic isolated subscapularis tendon lesions Lafosse III-IV: a prospective MRI-controlled case series with one year follow-up
-

Section 2. The Work Under Consideration for Publication

Did you or your institution at any time receive payment or services from a third party for any aspect of the submitted work (including but not limited to grants, data monitoring board, study design, manuscript preparation, statistical analysis, etc...)? No

Complete each item by typing an X in answer yes or not and completing the information requested if an answer is Yes. If you have more than one relationship, add lines.

1. Grant

☒ No ☐ Yes, money paid to you ☐ Yes, money paid to institution* Name of entity____
Comments†____

2. Consulting fee or honorarium

☒ No ☐ Yes, money paid to you ☐ Yes, money paid to institution* Name of entity____
Comments†____

3. Support for travel to meetings for the study or other purposes

☒ No ☐ Yes, money paid to you ☐ Yes, money paid to institution* Name of entity____
Comments†____

4. Fees for participation in review activities such as data monitoring boards, statistical analysis, end-point committees, and the like

☒ No ☐ Yes, money paid to you ☐ Yes, money paid to institution* Name of entity____
Comments†____

5. Payment for writing or reviewing the manuscript

☒ No ☐ Yes, money paid to you ☐ Yes, money paid to institution* Name of entity____
Comments†____

6. Provision of writing assistance, medicines, equipment, or administrative support

☒ No ☐ Yes, money paid to you ☐ Yes, money paid to institution* Name of entity____
Comments†____

7. Other

☒ No ☐ Yes, money paid to you ☐ Yes, money paid to institution* Name of entity____
Comments†____

* This means money that your institution received for your efforts on this study.

† Use this section to provide any needed explanation.

Section 3. Relevant financial activities outside the submitted work

1. Board membership

☐X___No ___Yes, money paid to you ___Yes, money paid to institution* Name of entity___ Comments___

2. Consultancy

☐X___No ___Yes, money paid to you ___Yes, money paid to institution* Name of entity___ Comments___

3. Employment

☐X___No ___Yes, money paid to you ___Yes, money paid to institution* Name of entity___ Comments I'm currently employed at the University Hospital Basel, Section Vascular Surgery.

4. Expert testimony

☐X___No ___Yes, money paid to you ___Yes, money paid to institution* Name of entity___ Comments___

5. Grants/grants pending

☐X___No ___Yes, money paid to you ___Yes, money paid to institution* Name of entity___ Comments___

6. Payment for lectures including service on speakers bureaus

☐X___No ___Yes, money paid to you ___Yes, money paid to institution* Name of entity___ Comments___

7. Payment for manuscript preparation

☐X___No ___Yes, money paid to you ___Yes, money paid to institution* Name of entity___ Comments___

8. Patents (planned, pending or issued)

☐X___No ___Yes, money paid to you ___Yes, money paid to institution* Name of entity___ Comments___

9. Royalties

☐X___No ___Yes, money paid to you ___Yes, money paid to institution* Name of entity___ Comments___

10. Payment for development of educational presentations

☐X___No ___Yes, money paid to you ___Yes, money paid to institution* Name of entity___ Comments___

11. Stock/stock options

☐X___No ___Yes, money paid to you ___Yes, money paid to institution* Name of entity___ Comments___

12. Travel/accommodations/ meeting expenses unrelated to activities listed**

☐X___No ___Yes, money paid to you ___Yes, money paid to institution* Name of entity___ Comments___

13. Other (err on the side of full disclosure)

☐X___No ___Yes, money paid to you ___Yes, money paid to institution* Name of entity___ Comments___

** For example, if you report a consultancy above there is no need to report travel related to that consultancy on this line.

Section 4. Other relationships

Are there other relationships or activities that readers could perceive to have influenced, or that give the appearance of potentially influencing, what you wrote in the submitted work?

 X No other relationships/conditions/circumstances that present a potential conflict of interest

___ Yes, the following relationships/conditions/circumstances are present (explain below):

>>>>>>>>>>><<<<<<<<<<<<

The International Committee of Medical Journal Editors

The ICMJE Disclosure of Potential Conflicts of Interest Form was adopted by *Arthroscopy: The Journal of Arthroscopic and Related Surgery* along with 17 other leading orthopaedic journals at the 2011 annual meeting of the American Academy of Orthopaedic Surgeons.